

OPEN ACCESS

Citation: Speed C, Arneil T, Harle R, Wilson A, Karthikesalingam A, McConnell M, et al. (2023) Measure by measure: Resting heart rate across the 24-hour cycle. PLOS Digit Health 2(4): e0000236. https://doi.org/10.1371/journal.pdig.0000236

Editor: Valentina Lichtner, University of Leeds, UNITED KINGDOM

Received: February 17, 2022 Accepted: March 20, 2023 Published: April 28, 2023

Copyright: © 2023 Speed et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: The study data can be made available on request. To request access to the de-identified data please email chrt-lon-data@google.com from an institutional email address. This is an email address that is independent of the authors (i.e. it is not an email address of any author) and this third party (not any of the authors) will provide data access. Please provide the following information: your name, position, the postal address of your institution and a contact telephone number. Please also provide a statement describing how you intend to use the data. Once approved, you will be provided with a

RESEARCH ARTICLE

Measure by measure: Resting heart rate across the 24-hour cycle

Cathy Speed 1,2,30 *, Thomas Arneil †, Robert Harle 1,40, Alex Wilson †, Alan Karthikesalingam †, Michael McConnell 1,50, Justin Phillips 1

- 1 Google, Mountain View, California, United States of America, 2 Cambridge Health & Performance, Cambridge, United Kingdom, 3 Cardiff Metropolitan University, Cardiff, United Kingdom, 4 Computer Laboratory, University of Cambridge, Cambridge, United Kingdom, 5 Division of Cardiovascular Medicine, Stanford University School of Medicine, Stanford, California, United States of America
- These authors contributed equally to this work.
- ‡ TA, AW and AK also contributed equally to this work.
- * speedc@google.com

Abstract

Background

Photoplethysmography (PPG) sensors, typically found in wrist-worn devices, can continuously monitor heart rate (HR) in large populations in real-world settings. Resting heart rate (RHR) is an important biomarker of morbidities and mortality, but no universally accepted definition nor measurement criteria exist. In this study, we provide a working definition of RHR and describe a method for accurate measurement of this biomarker, recorded using PPG derived from wristband measurement across the 24-hour cycle.

Methods

433 healthy subjects wore a wrist device that measured activity and HR for up to 3 months. HR during inactivity was recorded and the duration of inactivity needed for HR to stabilise was ascertained. We identified the lowest HR during each 24-hour cycle (true RHR) and examined the time of day or night this occurred. The variation of HR during inactivity through the 24-hour cycle was also assessed. The sample was also subdivided according to daily activity levels for subset analysis.

Findings

Adequate data was obtained for 19,242 days and 18,520 nights. HR stabilised in most subjects after 4 minutes of inactivity. Mean (SD) RHR for the sample was 54.5 (8.0) bpm (day) and 50.5 (7.6) bpm (night). RHR values were highest in the least active group (lowest MET quartile). A circadian variation of HR during inactivity was confirmed, with the lowest values being between 0300 and 0700 hours for most subjects.

Interpretation

RHR measured using a PPG-based wrist-worn device is significantly lower at night than in the day, and a circadian rhythm of HR during inactivity was confirmed. Since RHR is such link and instructions for downloading the data from a secure repository.

Funding: This study was entirely funded by Google, and all authors received employee salaries from Google during the research study. No additional grants or commercial funding sources were involved in this study. The funder had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing interests: I have read the journal's policy and the authors of this manuscript have the following competing interests: All authors are employees of Google, with the exception of TC Arneil who was an employee of Google at the time of the research study. All authors who are employees have stock options with Alphabet.

an important health metric, clarity on the definition and measurement methodology used is important. For most subjects, a minimum rest time of 4 minutes provides a reliable measurement of HR during inactivity and true RHR in a 24-hour cycle is best measured between 0300 and 0700 hours. Funding: This study was funded by Google.

Introduction

Resting heart rate (RHR) is an important biomarker of health and disease [1–7]. It is a vital sign, an indicator of all-cause mortality [1,2,7], cardiometabolic risk [1–9], cardiorespiratory fitness [10] and is used in regular monitoring to indicate general response to training [10] and is a highly variable physiological marker with many influencing factors. There is considerable inconsistency among definitions and methodological criteria for assessment of RHR, and consumer devices that provide estimations of RHR to their users show considerable variability in methodology and reported values. The term "resting heart rate" sometimes is used to represent the heart rate (HR) of an individual whenever they are inactive, as opposed to the lowest HR at rest in a 24-hour cycle. Methodological variables including time of measurement, subject positioning, pre-measurement rest, and mode of measurement (pulse palpation, ECG, and photoplethysmography (PPG) measured using wearable technologies such as a wristband or watch) may all influence the measurement. With the increasing availability of continuous HR monitoring using wearable devices, and the expansion of use of these devices in health monitoring, clarity on definitions and the differences between measurements is important, and within reach.

PPG-sensor technology has been demonstrated to accurately measure HR when compared to ECG and is increasingly used in cardiovascular health studies [11,12]. Note that PPG devices vary in their design and algorithms used to calculate HR in activity and at rest. The use of PPG-based wearable devices offers new opportunities in research across large populations. In a retrospective study of over 92,000 subjects wearing a range of different wrist-worn devices, mean 'RHR' was reported to range from 40–109 bpm across a US-based population with a mean (SD) age 45.8 (14.4) years [13]. RHR was provided using a proprietary formula from the device maker, with a single value being provided for each day, but no further insights were provided as to the methodology of its estimation. To our knowledge, no other PPG-based study has provided information as to methodology of RHR calculation.

In this real-world study, our primary aims were to describe a method for measurement of RHR recorded using PPG derived from wrist-worn fitness tracker measurements, to demonstrate the potential for erroneous estimation of RHR, and to assess quantitatively how long it takes HR to stabilise after activity ceases. We also evaluated the time of day on HR when inactive, examining for the presence of a circadian rhythm of HR during inactivity across 24 hours, and we evaluated the timing of RHR across subjects. We also further scrutinised the lowest HR during inactivity in the day compared to values at night.

Materials and methods

Study population

The study targeted healthy adults living within the United States aged 18–65 years. Eligible participants were physically active, as assessed by the International Physical Activity Questionnaire (IPAQ) [14,15]. The study was approved by an Institutional Review Board (Advarra, Columbia, MD). Written informed consent was obtained from each subject. Consenting

individuals agreed to wear a wrist device, the Fitbit Charge 4 fitness tracker (Fitbit LLC, San Francisco, CA, USA), continuously for up to three months. The study population comprised 433 subjects (66.1% female), mean(+-SD) age = 35.4 +- 9.6 years. Mean (+-SD) BMI was 26.5 (+- 5.2). No subjects had known cardiovascular disease and none was on antiarrhythmics. Subjects with a pacemaker and those who were night shift workers were excluded.

Data collection

The wrist-worn device captured HR using PPG sensors and activity through accelerometry. Participants were asked to perform their normal daily and night-time routines while wearing the device as often as possible. The device charges rapidly and subjects were asked to charge the device whilst bathing. Data from each device was uploaded automatically to a central secure platform (the Fitabase research platform [16]).

HR and estimates of activity level from accelerometry were recorded by the device every 60 seconds. HR during inactivity across the day and night was derived from these HR by taking the lowest HR measured by the device when still, i.e. when the metabolic equivalent (MET) during each 60 second epoch was estimated to be less than 1.5. The device itself has filtering algorithms to remove noise, although it should be noted that when subjects are inactive, the HR accuracy is typically reliable. Chow et al. found that the mean absolute percentage error (MAPE) of HR measured using fitness trackers in young adults at rest compared to an ECG chest strap (Polar M3) was 3.96% and 4.46% for the Garmin Vivosmart HR+ and Xiaomi Mi Band 2 devices respectively [17].

The wrist-worn device has built-in algorithms to determine the start and end times that the wearer was in bed, which were logged to enable separate calculations of lowest HR during inactivity in the day as well as at night. This implicitly defined night-time for a user as the primary sleep period identified by the tracker's algorithms.

Data analysis

Time series of HR and physical activity were analysed to produce estimates of daytime and night-time HR during inactivity. The RHR for each day and each night were determined by taking the lowest HR recorded when the wearer was inactive during the day and at night, respectively. Night-time and daytime heart rates were distinguished from automated logging of bed start and end times by the Fitbit device. For daytime calculations, the device must have been worn for at least 8 hours in the daytime and for night-time calculations, for at least 3 hours at night.

Activity recognition on the device allowed sedentary (MET < 1.5) times to be identified, and HR during those periods was scrutinised. The time taken for the HR to stabilise during inactivity was examined to identify the necessary duration of rest needed to allow best estimate of the RHR value.

HR over the daytime and night-time were compared and the variation of HR during inactivity across the 24-hour cycle was investigated. We also examined the effects of average activity levels on RHR.

Role of the funding source: The study was funded by Google, who funded all personnel and devices.

Results

Activity tracking confirmed the subjects to be physically active, with a mean (SD) number of minutes of moderate/vigorous physical activity 46.3 (28.5) minutes per day. The guidelines for

minimum levels of physical activity for health recommend at least 150 minutes of MVPA per week [18]. No significant change in activity levels was seen over the course of the study period.

Of 26,758 possible subject days and nights in the study, a total of 19,242 days of daytime data (>8 hours of use), and 18,520 nights (>3 hours of sleep) of data were obtained. This represents 71.9% compliance for daytime wear, and 69.2% compliance for night-time use. Of the qualifying data, the mean (SD) hours worn in a 24hr period was 15.19 (2.01) hours and at night was 7.97 (1.41) hours. The night-time mean start (bedtime) and finish (rise) times were 23:02 and 06:27, respectively.

To identify the minimum acceptable time for the HR to stabilise, periods of inactivity of 20 minutes or more were identified. This identified 160,357 sections of data (day) and 44,502 (night). The number of sampling sections was lower at night as the duration of each sampling period was longer when the subjects were consistently inactive when asleep. The HR recorded during each minute was compared with the HR from the preceding minute. The HR differences for adjacent minutes for the first 7 minutes during day and night are shown in Table 1. The mean decrease in HR between 3 and 4 minutes after a period of inactivity begins is less than 0.5 bpm. After 4 minutes of inactivity, a further mean decrease in HR of less than 1.0 bpm was observed when compared to 20 minutes of inactivity. The mean HR values recorded during each inactive minute are shown in Fig 1.

There are many examples in the data where the HR took longer to stabilise. This phenomenon was investigated by examining the percentage of the sample where HR during inactivity fell by more than 1, 2 or 3 beats per minute in each time interval. The results of this analysis are shown in Table 2. The HR in over 13% of subjects had not stabilised, i.e., the change in HR from 14 to 15 minutes of inactivity is greater than 3 bpm.

RHR in a 24-hour period was taken to be the lowest HR recorded during day or night after at least four minutes of inactivity. We also compared values of the lowest HR during inactivity in the day with that at night. Night-time values were lower than those in daytime, as shown in Table 3. Note the low level of 34 bpm was measured in one very fit subject on one occasion; the mean RHR for this subject during the study was 38 bpm. The mean (SD) difference (daynight) was + 3.9 (3.8) bpm. A paired Student's t-test showed significant differences between day and night-time RHR measurements (t-statistic = 136.99, P < 0.0001). The distribution of true RHR is illustrated in Fig 2.

The daytime lowest HR during inactivity obtained using our method was compared with the daytime 'RHR' calculated using the only other available definition, i.e.: the lowest HR during the day irrespective of activity [13]. The latter 'RHR' mean (SD) was 53.9 (9.9), i.e. 0.6 bpm lower than our estimate, due to a small number of instances where the lowest daily HR occurs when the subject is deemed to be moving by the device. The night-time lowest HR estimates differed by less than 0.1 bpm.

Table 1. Change in heart rate during inactivity.

	DAYTIME ($n = 160,357$)	NIGHTTIME $(n = 44,502)$		
Adjacent inactive mins	Mean decrease in HR from preceding minute (bpm)	Mean decrease in HR from preceding minute (bpm)		
0-1	2.99	1.63		
1-2	3.22	2.21		
2-3	0.96	0.41		
3-4	0.42	0.24		
4-5	0.24	0.14		
5-6	0.17	0.06		
6-7	0.16	0.03		

https://doi.org/10.1371/journal.pdig.0000236.t001

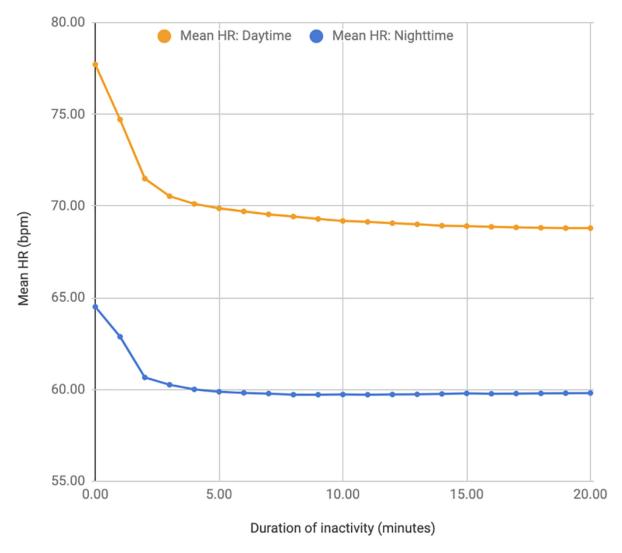


Fig 1. Decline in mean heart rate after initiation of inactivity.

https://doi.org/10.1371/journal.pdig.0000236.g001

We examined the trend in mean HR during inactivity across the 24-hour cycle during the study period (Fig 4). All HR measurements after at least 4 minutes of inactivity were averaged for all subjects for each hour of the day as indicated by the devices' timestamps.

Lastly, we examined the effects of individuals' activity levels on their lowest HR during inactivity (Table 4). The activity level for each subject was represented by the mean number of minutes of the day where the MET, estimated from the device's accelerometer, was greater than 3.0, equivalent to standard definitions of moderate to vigorous activity (MVPA). The results were grouped by activity level quartiles (Fig. 5).

Between-group differences in RHR were tested for significance using a single-factor ANOVA analysis with post-hoc subgroup comparisons. Table 4 shows the results from a multiple comparison of means using Tukey's honestly significant difference (HSD) test with a family-wise error rate (FWER), i.e. the probability of a Type I error, set to 0.05. The lowest activity and highest activity quartiles had mean (SD) RHRs of 55.2(7.94) bpm and 48.5(6.51) bpm respectively. We noted that those in the first activity quartile (i.e., individuals with lowest

Adjacent inactive mins	% of sample where change in HR > 1 bpm	% of sample where change in HR > 2 bpm	% of sample where change in HR > 3 bpm			
		DAYTIME				
0-1	0-1 54.9 46.9					
1-2	61.3	51.5	42.7			
2-3	44.2	33.0	24.4			
3-4	38.7	27.0	18.6			
14-15	33.0	21.1	13.1			
	N	NIGHTTIME				
0-1	48.9	40.8	34.3			
1-2	52.3	42.0	34.3			
2-3	35.9	23.4	15.9			
3-4	32.8	19.1	11.7			
14-15	23.8	12.7	7.6			

Table 2. Percentage of sample where heart rate during inactivity had not stabilised.

https://doi.org/10.1371/journal.pdig.0000236.t002

mean activity levels) had significantly higher RHR's compared to those in the other three quartiles.

Discussion

Several studies have demonstrated an association between RHR and cardiovascular mortality both in general populations [1–3] and in patients with hypertension [4,5] or cardiac diseases [6,7]. 'RHR' in the studies was taken as heart rate when inactive. HR during inactivity in this context has predominantly been studied during the day [1–7] and has variably been defined as a single measure on ECG [1], or pulse palpation [2,5,7]. Some studies that have demonstrated an association between resting heart rate and all-cause and cardiovascular disease mortality have empirically used daytime RHR after at least 5 minutes of supine rest, without providing a rationale for the rest period [19,20].

Night-time HR has been much less commonly studied but has been reported by some to be a better predictor of cardiovascular events than day-time lowest HR [21,22]. One study that examined participants where more than half of the individuals had hypertension failed to demonstrate daytime RHR association with mortality in contrast to the large body of evidence suggesting otherwise [23]. The use of wearable devices offers the opportunity for further examination of RHR across different times of the day and night and the relationship to cardiovascular disease and mortality in healthy and diseased populations.

It is notable that the HR in some subjects took some minutes to stabilise, and indeed in over 13% of subjects it had not fully stabilised (i.e., it was decreasing at a rate greater than 3 bpm) after 15 minutes of inactivity. These results were considered most likely to have occurred due to the physiological effect of more intense exercise before the rest period, and the finding that, in most subjects the HR had stabilised after 4 minutes, we suggest that in most subjects a minimum of 4 minutes rest is acceptable, but the activity level of the subject in the few hours

Table 3. Summary of daytime and night-time lowest heart rates during inactivity.

	N	Mean(+-SD) Heart rate	Heart rate Range		
Daytime	19242	54.5 (8.0)	34.0-92.4		
Nighttime	18520	50.5 (7.6)	32.6-89.8		

https://doi.org/10.1371/journal.pdig.0000236.t003

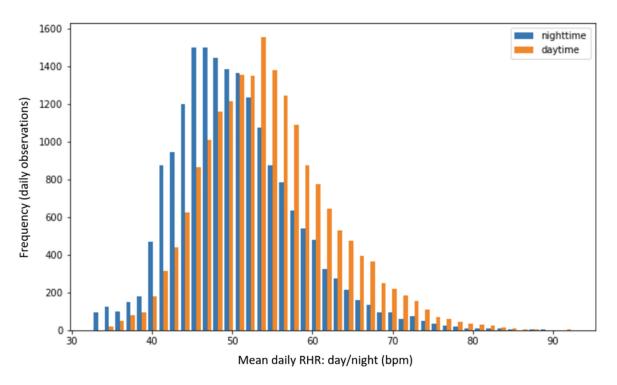


Fig 2. The distribution of mean RHR estimates for both daytime and night-time. We also examined the time of day that lowest HR was noted. Just over half (53%) occurred between 0300 and 0700. The distribution is shown in the histogram below (Fig 3).

https://doi.org/10.1371/journal.pdig.0000236.g002

prior to measurement should be considered. This has relevance for clinical practice, and studies where "resting heart rate" is taken by pulse after only a few minutes of rest.

The use of PPG in the monitoring of HR allows large-scale, real-world population studies that are not feasible using ECG. The increasing quality of PPG sensors has resulted in excellent quality of HR measurement in newer devices such as that used in this study. At rest, their accuracy is similar to ECG [11,12,24,25], and the inclusion of activity recognition in our study allowed us to ascertain when the subject was inactive. This is important, as when there is error in signal detection with wrist-worn devices, this is more likely when the subject is active, especially at vigorous intensity, and erroneously low HR during such activity can occur. 'Ambulatory' HR has also been proposed to be relevant, and also has varying definitions, from the mean derived from 24-hour continuous monitoring, or from 'snapshots' measured during the day. It too has also been reported to have predictive value in mortality risk in elderly hypertensives [26]. Whilst it is possible to measure this using PPG, there is more likely to be error due to activity affecting signal integrity, and the capacity to measure true RHR may make 'ambulatory' recordings unnecessary for risk assessment.

As anticipated, RHR in our study was noted to be significantly lower in those subjects who were routinely more active, but a 'little goes a long way', with the most significant difference in RHR being seen between the least active and the next quartile of activity level.

We noted a circadian rhythm of HR during inactivity, and this has been studied by a few groups, who have also noted the trend we identified in our study, specifically with lower HR during the night and in the first few hours in the morning [27,28]. This is considered to be related to the dominance of sympathetic tone and circulating catecholamines during the day, and its relative decline at night, accompanied by increased parasympathetic tone during those

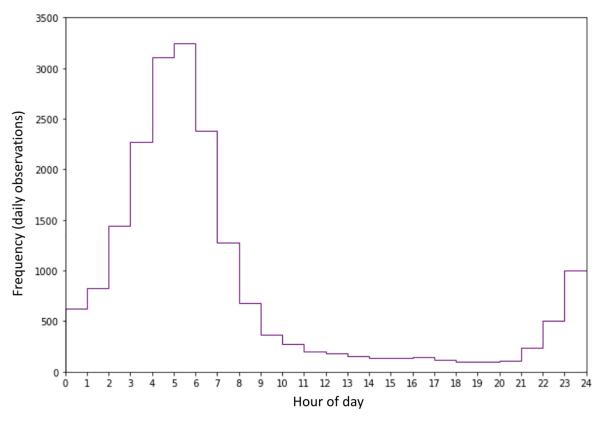


Fig 3. Distribution across subjects of timing of lowest RHR in 24-hour cycle, showing that on most days the RHR was lowest between 0300 and 0700 across the population.

https://doi.org/10.1371/journal.pdig.0000236.g003

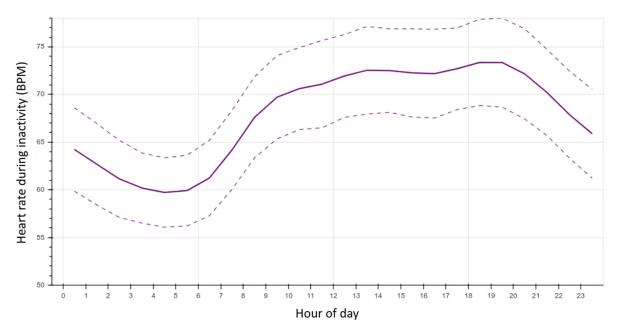


Fig 4. Circadian variation in heart rate during inactivity. Mean (SD) heart rate after 4 minutes of inactivity by hour of the day.

https://doi.org/10.1371/journal.pdig.0000236.g004

Table 4.	Multiple	Comparisor	ı of Means-	—Tukey	HSD	(FWER = 0.05).
----------	----------	------------	-------------	--------	-----	----------------

Group (1)	Group (2)	Mean diff (bpm)	p-adj	Lower	Upper	Reject?
1st	2nd	-3.7005	0.0019	-6.3432	-1.0578	True
1st	3rd	-4.0132	0.001	-6.6559	-1.3705	True
1st	4th	-6.7866	0.001	-9.4293	-4.1439	True
2nd	3rd	-0.3127	0.900	-2.9554	2.330	False
2nd	4th	-3.0861	0.0146	-5.7288	-0.4434	True
3rd	4th	-2.7734	0.0355	-5.4161	-0.1307	True

https://doi.org/10.1371/journal.pdig.0000236.t004

night-time hours [27]. Activity between rest intervals may also influence the values across the day, and this may have influenced our finding that in some samples the HR can take longer than the typical 4 minutes to fully stabilise.

It can be proposed from this data that true RHRs occur most commonly between 0300 and 0700 in the general population, apart from those who are night shift workers, who were excluded from this study.

A limitation of this study is that it may not be wholly generalizable to other wearable devices. Our estimates of HR were dependent on the Fitbit Charge 4's segmentation of day/ night and moving/sedentary periods. These factors may introduce some bias in our recommended approach to measuring RHR, so we suggest a comparison of results from a range of wearable devices as an area for future research.

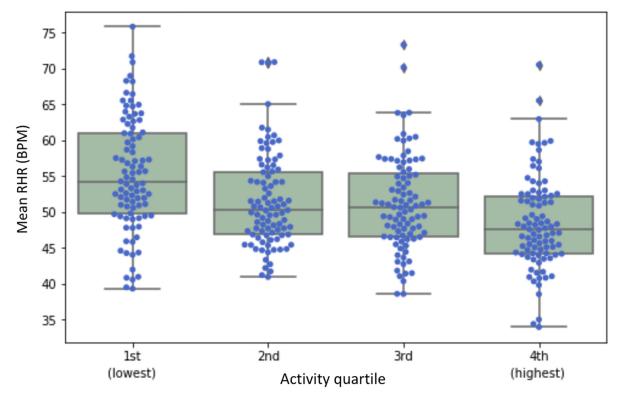


Fig 5. Box plots of the mean RHR in the 24-hour cycle for each activity quartile.

https://doi.org/10.1371/journal.pdig.0000236.g005

Conclusions

Resting heart rate is a vital physiological marker that reflects health, disease and mortality risk. Nevertheless, specific guidance as to how to measure it reliably is lacking. Before the availability of reliable wrist-worn devices using PPG signals, measurement has relied largely on pulse palpation and ECG-based measures. The use of PPG-based devices such as those used in our study allows specific criteria to be set relating to RHR measurement across all settings, and greater insights into its circadian variation. We recommend a minimum inactivity period of 4 minutes for reliable HR measurement to calculate RHR and suggest that subjects have also not engaged in significant exercise in the immediately preceding period. We also recommend that the true RHR in a 24-hour cycle is best measured between 0300 and 0700 hours. We encourage authors to clarify their methodology of RHR measurement in future studies.

Author Contributions

Conceptualization: Cathy Speed.

Data curation: Justin Phillips.

Formal analysis: Cathy Speed, Thomas Arneil, Robert Harle, Alex Wilson, Alan Karthikesalingam, Michael McConnell, Justin Phillips.

Investigation: Cathy Speed, Robert Harle, Justin Phillips.

Methodology: Cathy Speed, Robert Harle, Alex Wilson, Michael McConnell, Justin Phillips.

Supervision: Robert Harle, Michael McConnell.

Writing – original draft: Cathy Speed, Robert Harle, Alex Wilson, Alan Karthikesalingam, Michael McConnell, Justin Phillips.

Writing - review & editing: Cathy Speed, Robert Harle, Michael McConnell, Justin Phillips.

References

- Kannel WB, Kannel C, Paffenbarger RS Jr, Cupples LA. Heart rate and cardiovascular mortality: The Framingham Study. Am Heart J. 1987; 113:1489–1494. https://doi.org/10.1016/0002-8703(87)90666-1 1 PMID: 3591616
- Palatini P, Casiglia E, Julius S, Pessina AC. High heart rate: a risk factor for cardiovascular death in elderly men. Arch Intern Med. 1999; 159:585–592. https://doi.org/10.1001/archinte.159.6.585 PMID: 10090115
- 3. Jouven X, Empana JP, Schwartz PJ, Desnos M, Courbon D, Ducimetiere P. Heart-rate profile exercise as a predictor of sudden death. N Engl J Med. 2005; 352:1951–1958.
- Kolloch R, Legler UF, Champion A, Cooper-Dehoff RM, Handberg E, Zhou Q et al. Impact of resting heart rate on outcomes in hypertensive patients with coronary artery disease: findings from the INternational VErapamil-SR/trandolapril STudy (INVEST). Eur Heart J. 2008; 29:1327–1334. https://doi.org/ 10.1093/eurheartj/ehn123 PMID: 18375982
- Paul L, Hastie CE, Li WS, Harrow C, Muir S, Connell JMC, et al. Resting Heart Rate Pattern During Follow-Up and Mortality in Hypertensive Patients. Hypertension. 2010; 55:567–574. https://doi.org/10.1161/HYPERTENSIONAHA.109.144808 PMID: 20038750
- Copie X, Hnatkova K, Staunton A, Fei L, Camm AJ, Malik M. Predictive power of increased heart rate versus depressed left ventricular ejection fraction and heart rate variability for risk stratification after myocardial infarction. Results of a two-year follow-up study. J Am Coll Cardiol. 1996; 27:270–276. https://doi.org/10.1016/0735-1097(95)00454-8 PMID: 8557893
- Diaz A, Bourassa MG, Guertin MC, Tardif JC. Long-term prognostic value of resting heart rate in patients with suspected or proven coronary artery disease. Eur Heart J. 2005; 26:967–974. https://doi. org/10.1093/eurheartj/ehi190 PMID: 15774493
- Caetano J. & Delgado Alves J. Heart rate and cardiovascular protection. Eur. J. Intern. Med. 2015; 26, 217–222.

- Alhalabi L, Singleton MJ, Oseni AO, Shah AJ, Zhang Z-M, Soliman EZ. Relation of Higher Resting Heart Rate to Risk of Cardiovascular Versus Noncardiovascular Death. The American Journal of Cardiology. 2017; 119, 1003–1007. https://doi.org/10.1016/j.amjcard.2016.11.059 PMID: 28132682
- American College of Sports Medicine. ACSM Guidelines for exercise testing and prescription. Tenth Edition. Philadelphia (PA): Wolters Kluwer, 2018.
- Wallen MP, Gomersall SR, Keating SE, Wisloff U, Coombes JS. Accuracy of Heart Rate Watches: Implications for Weight Management. PloS one. 2016; 11(5):e0154420. https://doi.org/10.1371/journal.pone.0154420 PMID: 27232714
- Shcherbina A, Mattsson CM, Waggott D, Salisbury H, Christle JW, Hastie T, et al. Accuracy in Wrist-Worn, Sensor-Based Measurements of Heart Rate and Energy Expenditure in a Diverse Cohort. J Pers Med. 2017; 7(2):3. https://doi.org/10.3390/jpm7020003 PMID: 28538708
- 13. Quer G, Gouda P, Galarnyk M, Topol EJ, Steinhubl SR. Inter- and intraindividual variability in daily resting heart rate and its associations with age, sex, sleep, BMI, and time of year: Retrospective, longitudinal cohort study of 92,457 adults. PLOS ONE 2020; 15(2): e0227709. https://doi.org/10.1371/journal.pone.0227709 PMID: 32023264
- Booth M.L. Assessment of Physical Activity: An International Perspective. Research Quarterly for Exercise and Sport, 2000; 71 (2): s114–20.
- 15. https://www.ipaq.ki.se.
- 16. https://www.fitabase.com.
- Chow H, and Chao-Ching Y. Accuracy of Optical Heart Rate Sensing Technology in Wearable Fitness Trackers for Young and Older Adults: Validation and Comparison Study." JMIR mHealth and uHealth 2020; 8(4): e14707. https://doi.org/10.2196/14707 PMID: 32343255
- 18. https://www.cdc.gov/physicalactivity/basics/adults/index.html.
- Kristal-Boneh E. Silber H, Harari G, Froom (2000) 'The association of resting heart rate with cardiovascular, cancer and all-cause mortality. Eight year follow-up of 3527 male Israeli employees (the CORDIS Study)', European heart journal, 21(2), pp. 116–124. https://doi.org/10.1053/euhj.1999.1741 PMID: 10637085
- Benetos A, Rudnichi A, Thomas F, Safar M, Guize L. Influence of heart rate on mortality in a French population: role of age, gender, and blood pressure. Hypertension. 1999 Jan; 33(1):44–52. https://doi. org/10.1161/01.hyp.33.1.44 PMID: 9931080
- Palatini P, Reboldi G, Beilin LJ, Eguchi K, Imai Y, Kario K, et al. Predictive Value Of Night-time Heart Rate For Cardiovascular Events In Hypertension. The Abp International Study. Int J Cardiol. 2013; 168 (2): 1490–5.
- Johansen CD, Olsen RH, Pedersen LR, Kumarathurai P, Mouridsen MR, Binici Z et al. Resting, nighttime, and 24 h heart rate as markers of cardiovascular risk in middle-aged and elderly men and women with no apparent heart disease, *European Heart Journal*, Volume 34, Issue 23, 14 June 2013, Pages 1732–1739, https://doi.org/10.1093/eurhearti/ehs449 PMID: 23306958
- 23. Ben-Dov IZ, Kark JD, Ben-Ishay D, Mekler J, Ben-Arie L, Bursztyn M. Blunted heart rate dip during sleep and all-cause mortality. Arch Intern Med. 2007; 167:2116–2121. https://doi.org/10.1001/archinte. 167.19.2116 PMID: 17954807
- 24. Haghayegh S, Khoshnevis S, Smolensky MH, Diller KR. Accuracy of PurePulse photoplethysmography technology of Fitbit Charge 2 for assessment of heart rate during sleep. Chronobiology International 2019; 36: 927–933. https://doi.org/10.1080/07420528.2019.1596947 PMID: 30990098
- **25.** Heneghan C, Venkatraman S, and Russell A. Investigation of an estimate of daily resting heart rate using a consumer wearable device. medrxiv 2019;19008771.
- Palatini P, Thijs L, Staessen JA, Fagard RH, Bulpitt CJ, Clement DL, et al. Predictive Value of Clinic and Ambulatory Heart Rate for Mortality in Elderly Subjects With Systolic Hypertension. Arch Intern Med. 2002; 162(20):2313–2321 https://doi.org/10.1001/archinte.162.20.2313 PMID: 12418945
- Scheer FA, van Doornen LJ, Buijs RM. Light and diurnal cycle affect human heart rate: possible role for the circadian pacemaker. J Biol Rhythms. 1999; 14(3):202–12. https://doi.org/10.1177/ 074873099129000614 PMID: 10452332
- 28. Malpas SC, Purdie GL. Circadian variation of heart rate variability. Cardiovasc Res. 1990; 24(3):210–3. https://doi.org/10.1093/cvr/24.3.210 PMID: 2346954